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ubstitute	o for form 1449A/PTO			Complete if Known			
	INFORMATION DISCLOSURE		Application Number	10/758,242			
1			Filing Date	January 16, 2004			
	STATEMENT BY API	PLIC	ANT	First Named Inventor	Bernd SUNDERMANN		
	•			Art Unit	1621		
(use as many sheets as necessary)		Examiner Name	Unassigned				
heet	1	of	5	Attorney Docket Number	029310.53093US		

U.S. PATENT DOCUMENTS								
Examiner	Cite	Document Number	Publication Date	Name of Patentee or	Pages, Columns, Lines, Where			
Initials'	No.1	Number-Kind Code <sup>2</sup> (if known)	MM-DD-YYYY	Applicant of Cited Document	Relevant Passages or Relevant Figures Appear			
BJD	AA	US- 5,304,479	04-19-1994	Cheng-I Lin	-			
BOD	AB	US- 5,239,110	08-24-1993	John P. Mallamo et al.				
Bab	AC	US- 4,366,172	12-28-1982	Daniel Lednicer				
BJO	AD	US- 4,346,101	08-24-1982	Daniel Lednicer				
BJb	AE	US- 4,212,878	07-15-1980	Daniel Lednicer et al.				
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	FOREIGN PATENT DOCUMENTS							
		Foreign Patent Document			Pages, Columns, Lines, Where Relevant Passages			
Examiner Initials'	Cite No.1	Country Code <sup>3</sup> -Number <sup>4</sup> -Kind Code <sup>6</sup> (if known)	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	or Relevant Figures Appear	τ°		
Bab	AG	DE 2839891	04-12-1979	The Upjohn Co.		AB		
RJO	AH	DE 19963175	07-12-2001	Gruenenthal GmbH	,	AB		
B 500	Al	WO 01/12195	02-22-2001	Gruenenthal GmbH		AB		
BJO	AJ	EP 0410191	01-30-1991	Bayer AG		AB		

		NON PATENT LITERATURE DOCUME	NTS				
Examiner Initials*	I include name of the author (in CAPITALLETTERS), title of the article (when appropriate), title of the item (book						
Bob	AK	DANIEL LEDNICER ET AL., "4-(p-Bromophenyl)-4-(di phenethylcyclohexanol, an Extremely Potent Represer Journal of Medicinal Chemistry, October 1979, pp. 119 Chemical Society	ntative of a New Analgesic Series",				
BOD	AL	HIROSHI KAWAMOTO ET AL., "Synthesis of J-113 ORL1 Antagonist," Tetrahedron, 2001, pp. 981-986, 5					
	_ <b>A</b> M	PHILLIP F. VONVOIGTLANDER ET AL., "4-Aryl-4-al Chemical Nevel Series of Analgesics Including Opioi Agenist," pp. 17-21					
Examiner Signature	T	<i>K</i>	Date 12/27/11S				

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Substitute for	or form 1449B/PTO			Application Number	10/758,242	
1	NFORMATION DI	SCLOSUR	E	Filing Date	January 16, 2004	
STATEMENT BY APPLICANT				First Named Inventor	Bernd SUNDERMANN	
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Sheet	2	of	5	Attorney Docket Number	029310.53093US	

		NON PATENT LITERATURE DOCUMENTS	S				
Examiner Initials*	Cite No. <sup>1</sup>	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.					
Bod	AN	AUD A. ABDULLA ET AL., "Axotomy Reduces the Effect of Analgesic Opioids Yet ncreases the Effect of Nociceptin on Dorsal Root Ganglion Neurons," The Journal of Neuroscience, December 1, 1998, pp. 9685-9694, 18, 23, Society for Neuroscience					
bsp	AO	GIROLAMO CALO ET AL., "Pharmacology of Nocice Therapeutic Target," British Journal of Pharmacolo Macmillan Publishers Ltd.					
BJD	AP	MARK CONNER ET AL., "The Effect of Nociceptin Intracellular Ca <sup>2+</sup> in the SH-SY5Y Human Neuroblastor 118, Stockton Press					
820	AQ	E.S.L. FABER ET AL., "Depression of Glutamatergic To Neonatal Rat Hemisected Spinal Cord Preparation In V 1996, pp. 1-2,	ransmission litro", Specia	by Nociceptin in the al Report, July 19,			
BJD	AR	"Opioid and Opiate Receptors: Peptides and Knock-Ou 1998, p. 1358, Vol. 24	it," Society f	or Neuroscience,			
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g Z b	AU	TOSHIYA MANABE ET AL., "Facilitation of Long-Term Lacking Nociceptin Receptors", Letters To Nature, Aug 394, Macmillan Publishers Ltd.	Potentiation just 6, 1998,	n and Memory in Mice pp. 577-581, Vol.			
800	AV	JEAN-CLAUDE MEUNIER ET AL., "Isolation and Structure of Opiod Receptor-Like ORL, Receptor," Letters to Nat 535, Vol. 377					
Examiner		6/	Date	12/27/05			

Considered Signature

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Substitute to	10m 1449B/P10			Application Number 10/758,242			
41	FORMATION E	DISCLOSUE	?F	Filing Date	January 16, 2004		
	TATEMENT BY			First Named Inventor	Bernd SUNDERMANN	SUNDERMANN	
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Sheet	3	of	5	Attorney Docket Number	029310.53093US		

		NON PATENT LITERATURE DOCUMENT	S						
Examiner Initials*	Cite No. <sup>1</sup>	Include name of the author (in CAPITAL LETTERS), title of the article magazine, journal, serial, symposium, catalog, etc.), date, page(s), vol country where published.			T2				
BJD	AW		J.S. MOGIL ET AL., "Orphanin FQ is a Functional Anti-Opioid Peptide", Neuroscience, 1996, pp. 333-337, Vol. 75, No. 2, Elsevier Science Ltd., Great Britain						
BJO	AX	Hearing Ability in Mice Lacking the Nociceptin/Orp	IIYUKI NISHI ET AL., "Unrestrained Nociceptive Response and Disregulation of learing Ability in Mice Lacking the Nociceptin/OrphaninFQ Receptor," The EMBO ournal, 1997, pp. 1858-1864, Vol. 16, No. 8, Oxford University Press						
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BJO	BA	TATSUO YAMAMOTO ET AL., "Effects of Intrathecally Administered Nociceptin, an Opioid Receptor-like, Receptor Agonist, and N-methyl-D-aspartate Receptor Antagonist on the Thermal Hyperalgesia Induced by Partial Sciatic Nerve Injury in the Rat," Anesthesiology, 1997, pp. 1145-1152, Vol. 87, No. 5, Lippincott-Raven Publishers							
aC8	ВВ	ALI ARDATI ET AL., "Interaction of [3 H]Orphanin FQ at the Orphanin FQ Receptor: Kinetics and Modulation by Nucleotides," Molecular Pharmacology, 1997, pp. 816-for Pharmacology and Experimental Therapeutics	Cations and	I Guanine					
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B30	BD	TRISTAN DARLAND ET AL., "Orphanin FQ/nociceptin But So Much More," TINS, 1998, PP. 215-221, Vol. 21							
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Sheet	4	of	5	Attorney Docket Number	029310.53093US	

		NON PATENT LITERATURE DOCUMENTS	
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BJD	BE	BULENT GUMUSEL ET AL., "Nociceptin: An Endogenous Agonist for Central Opioid Like <sub>1</sub> (ORL <sub>1</sub> ) Receptors Possesses Systemic Vasorelaxant Properties," Life Sciences, 1997, pp. PL 141-145, Vol. 60, No. 8, Elsevier Science Inc., USA	
B 55	BF	NAOKI HARA ET AL., "Characterization of Nociceptin Hyperalgesia and Allodynia in Conscious Mice," British Journal of Pharmacology, 1997, pp. 401-408, 121, Stockton Press	
BZD	BG	DANIEL R. KAPUSTA ET AL., "Diuretic and Antinatriuretic Responses Produced by the Endogenous Opioid-Like Peptide, Nociceptin (Orphanin FQ)," Life Sciences, 1997, pp. PL 15-21, Vol. 60, No. 1, Elsevier Science Inc., USA	
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BJD	ВІ	HANS MATTHES ET AL., "Functional Selectivity of Orphanin FQ for Its Receptor Coexpressed with Potassium Channel Subunits in Xenopus <i>laevis</i> Oocytes," Molecular Pharmacology, 1996, pp. 447-450, 50, The American Society for Pharmacology and Experimental Therapeutics	
BIP	BJ	JEFFREY S. MOGIL ET AL., "Functional Antagonism of $\mu$ -, $\delta$ - and $\kappa$ -opioid Antinociception by Orphanin FQ," Neuroscience Letters, 1996, pp. 131-134, 214, Elsevier Science Ireland Ltd.	
Bap	ВК	CATHERINE MOLLEREAU ET AL., "ORL1, A Novel Members of the Opioids Receptor Family Cloning, Functional Expression and Localization," FEBS Letters, 1994, 341, Federation of European Biochemical Societies	
350	BL	JAMES D. POMONIS ET AL., "Orphanin FQ, Agonist of Orphan Opioid Receptor ORL <sub>1</sub> , Stimulates Feeding in Rats," NeuroReport, December 20, 1996, pp. 369-371, Vol. 8, No.1, Rapid Science Publishers	

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B JD	ВМ	YS. SHU ET AL., "Orphanin FQ/Nociceptin Modulates Glutamate- and Kainic Acid- Induced Currents in Acutely Isolated Rat Spinal Dorsal Horn Neurons," Neuropeptides, 1998, pp. 567-571, 32, Harcourt Brace & Co., Ltd.	
BJD	BN	XIAO-JUN XU ET AL., "Nociceptin or Antinociceptin: Potent Spinal Antinociceptive Effect of Orphanin FQ/ Nociceptin in the Rat," NeuroReport, September 2 1996, Vol. 17, No. 13, Rapid Science Publishers	
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RUD	BP	M.N.A. RAO ET AL., "Quantitative Correlation Between Hydrophobicity and Analgesis Activity of 4-Amino 4-Arylcyclohexanols," Indian Drugs, 1985, pp. 252-257, 22, 5	
BJb	BQ	JEAN-MARC KAMENKA ET AL., "Orientation Structurale et Conformationnelle de la Fixation de la Phencyclidine dans le SNC," Eur. J. Med. Chem. 1984, pp. 255-260, 19, 3	
Boo	BR	DANIEL LEDNICER ET AL., "4-Amino-4-arylcyclohexanones and Their Derivatives, a Novel Class of Analgestics", J. Med. Chem., 1980, pp. 424-430, 23	
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